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Amendments to the Claims:

- 1. (currently amended) A Piroplasmid protein comprising, characterised in that said protein comprises an amino acid sequence having a similarity of at least 70% with the amino acid sequence depicted in selected from the group consisting of:
 - i) SEQ ID NO: 2 ef,
 - ii) immunogenic fragments of SEQ ID NO: 2,
 - iii) SEQ ID NO: 4,
 - iv) immunogenic fragments of SEQ ID NO: 4,
 - <u>v)</u> <u>SEQ ID NO: 6,</u>
 - vi) immunogenic fragments of SEQ ID NO: 6,
 - vii) SEQ ID NO: 8,
 - viii) immunogenic fragments of SEQ ID NO: 8,
 - ix) SEQ ID NO: 10, [[or an]] and
 - <u>x</u>) immunogenic fragment of <u>SEQ ID NO</u>: 10;

wherein said protein inhibits invasion of an organism of the family Piroplasmida.

- 2. -3. (cancelled)
- 4. (currently amended) <u>A nucleic</u> Nucleic acid, characterised in that wherein said nucleic acid encodes a protein according to claim 1, or an immunogenic fragment of said protein.
- 5.-6. (cancelled)
- 7. (currently amended) A cDNA fragment comprising a nucleic acid according to claim 4 one or more of the claims 4 6.
- 8. (currently amended) A recombinant Recombinant DNA molecule comprising
 - i) a nucleic acid according to one or more of the claims 4 6 claim 4, or

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<u>ii)</u> a cDNA fragment according to claim 7, said nucleic acid or said cDNA fragment being comprising (i);

wherein said cDNA fragment or said nucleic acid is under the control of a functionally linked promoter.

- 9. (currently amended) A live Live recombinant carrier comprising
 - i) a nucleic acid according to one or more of the claims 4 6 claim 4,
- <u>ii)</u> a cDNA fragment according to claim 7, said nucleic acid or said cDNA fragment being under the control of a functionally linked promoter comprising (i),
 - iii) a recombinant DNA molecule according to claim 8 comprising (i), or
 - <u>iv)</u> a recombinant DNA molecule comprising (ii);

wherein said cDNA fragment or said nucleic acid is under the control of a functionally linked promoter.

- 10. (currently amended) A host Host cell comprising
 - i) a nucleic acid according to one or more of the claims 4 6 claim 4,
- ii) a cDNA fragment according to claim 7 comprising (i), said nucleic acid or said eDNA fragment being under the control of a functionally linked promoter,
 - iii) a recombinant DNA molecule comprising (i) according to claim 8,
 - iv) a recombinant DNA molecule comprising (ii),
 - <u>v</u>) or a live recombinant carrier comprising (i),
 - vi) a live recombinant carrier comprising (ii),
 - vii) a live recombinant carrier comprising (iii), or
 - viii) a live recombinant carrier comprising (iv) according to claim 9;

wherein said nucleic acid or said cDNA fragment are under the control of a functionally linked promoter.

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11. (currently amended) A vaccine Vaccine comprising a protein according to claim 1 one or more of the claims 1 - 3 or an immunogenic fragment of said protein, a nucleic acid according to one or more of the claims 4-6, a cDNA fragment according to claim 7, a recombinant DNA molecule according to claim 8, a live recombinant carrier according to claim 9, or a host cell according to claim 10, or a combination thereof, and a pharmaceutically acceptable carrier.

(currently amended) The vaccine Vaccine according to claim 11, characterised in that 12. said vaccine comprises further comprising an adjuvant.

13. (currently amended) The vaccine Vaccine according to claim 11, one or more of the claims 11 - 12, characterised in that said vaccine comprises further comprising an additional immunoactive component or a nucleic acid encoding said additional immunoactive component.

(currently amended) A vaccine comprising Vaccine, characterised in that said vaccine 14. comprises an antibody against a protein according to claim 1 one or more of the claims 1 3 or an antibody against an immunogenic fragment of said protein, or a combination thereof, and a pharmaceutically acceptable carrier.

15. (cancelled)

- 16. (currently amended) A Use of a protein according to one or more of the claims 1-3 or an immunogenic fragment of said protein for the manufacture of a vaccine for prophylactic or therapeutic treatment of an infection or its clinical signs caused by a Piroplasmid organism comprising administering a vaccine comprising a protein according to claim 1.
- (currently amended) A Use of a nucleic acid sequence according to one or more of the 17. claims 4 - 6, a cDNA fragment according to claim 7, a recombinant DNA molecule according to claim 8, a live recombinant carrier according to claim 9, or a host cell according to claim 10 for

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the manufacture of a vaccine for prophylactic or therapeutic treatment of an infection or its clinical signs caused by a Piroplasmid organism comprising administering a vaccine comprising

a nucleic acid according to claim 4.

18. (currently amended) A diagnostic Diagnostic test for the detection of a nucleic acid

associated with a Piroplasmid organism comprising

, characterised in that the test comprises a nucleic acid, said nucleic acid being at (i)

least 70 % similar to the a nucleic acid sequence depicted in SEQ ID NO: 1, 3, 5, 7, or 9, or

(ii) a nucleic acid that is complementary to (i) said nucleic acid, or

(iii) a nucleic acid that hybridises to (i) under stringent conditions:

wherein either of the nucleic acids (i), (ii) and (iii) each have a length of at least 15 nucleotides.

19. (currently amended) A diagnostic Diagnostic test for the detection of antibodies against a

Piroplasmid organism comprising, characterised in that said test comprises a protein according

to claim 1 one or more of the claims 1 - 3, or an immunogenic fragment of said protein, or a

combination thereof.

20. (currently amended) A diagnostic Diagnostic test for the detection of antigenic material

from a Piroplasmid organism comprising, characterised in that said test comprises an antibody

against a protein according to claim 1 one or more of the claims 1 - 3 or an antibody against an

immunogenic fragment of said protein, or a combination thereof.

21. (new) A vaccine comprising a nucleic acid according to claim 4 and a pharmaceutically

acceptable carrier.

(new) A vaccine comprising a cDNA fragment according to claim 7 and a 22.

pharmaceutically acceptable carrier.

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23. (new) A vaccine comprising a recombinant DNA molecule according to claim 8 and a pharmaceutically acceptable carrier.

24. (new) A vaccine comprising a live recombinant carrier according to claim 9 and a pharmaceutically acceptable carrier.

25. (new) A vaccine comprising a host cell according to claim 10 and a pharmaceutically acceptable carrier.